Global Institute Of Health Science™

CERTIFICATE · PG DIPLOMA · FAST-TRACK PG DIPLOMA

CLINICAL RESEARCH & PHARMACOVIGILANCE

A rare combination in one programme — and one fee.

NEW AI MODULE | LIFETIME 1-TO-1 EXPERT | FREE CAREER COACHING ₹9,999 | 2 INTERNATIONAL CERTIFICATES | 100% ONLINE

NEW · AI MODULE **NOW WITH AI IN CLINICAL RESEARCH, PHARMACOVIGILANCE & DRUG SAFETY**

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Trusted by Healthcare Professionals Globally

Learn from global perspectives and prepare for impactful roles in hospitals, public health systems and healthcare organisations around the world.

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15+ years of experience **2,000+** placed globally **Lifetime** free expert tutoring

ACCREDITED & CERTIFIED

- **ISO 21001:2018** Educational organisation management
- **UASL** Germany / UK — quality management
- **GSAAA** USA — global accreditation
- **IPR, Govt. of India** Copyright approved

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Recorded candidate satisfaction to date

PROGRAMME OVERVIEW

Build Future-Ready Skills Across the Clinical Trial & Drug-Safety Lifecycle

Practical | Job-Oriented | ICH-GCP | Clinical Trials | Drug Safety | Signal Detection | MedDRA | AI Literacy

The GIHS Clinical Research & Pharmacovigilance programme brings together two closely connected professional domains: the conduct and management of clinical trials, and the monitoring of medicine safety. A trial generates safety information; pharmacovigilance provides the structured processes for evaluating and acting on it. Learning both gives you the whole arc, from first-in-human study to post-marketing surveillance.

The syllabus strengthens knowledge of ICH-GCP, trial design and operations, regulatory requirements, study documentation, site management and clinical monitoring — then moves into adverse-event reporting, case processing, signal detection, risk management, MedDRA and pharmacovigilance quality systems.

> **"We Teach What the Job Requires."**
>
> GIHS connects professional theory with workplace-oriented clinical-research and drug-safety workflows, so learners build practical competency in addition to their existing academic qualifications.

Why This Programme Stands Out

- Clinical Research + Pharmacovigilance in one integrated professional programme
- ICH-GCP, Schedule Y, ICMR and Indian GCP concepts
- Trial design, protocol / CRF, essential documents and informed consent
- Site selection, initiation, recruitment, monitoring and close-out
- QC, compliance, auditing and 21 CFR Part 11 awareness
- Adverse event and ADR evaluation and drug-safety reporting
- Single-case processing and case narrative writing
- PV databases, signal detection, risk assessment and risk management
- PSUR / PBRER, MedDRA, PV laws, SOPs, audits and inspections
- NEW dedicated AI module spanning clinical trials and pharmacovigilance
- Certificate, PG Diploma and Fast-Track learning pathways

The Programme at a Glance

10	3 / 4 / 12	100%	Lifetime	MCQ + Project	2
Modules, including the new AI module	Months — Certificate, Fast Track or PG Diploma	Online & distance, study at your own pace	Free personal expert assigned to you	Online exam plus a 30-page project	International certificates awarded and shipped

REVISED SYLLABUS

Curriculum, Module by Module

The full GIHS syllabus, regrouped into 10 clearly sequenced modules so you can see exactly what each stage of the programme teaches and why it comes where it does. Module 10 is the new Artificial Intelligence module.

MODULE 01 Foundations of Clinical Research

What clinical research is, how it differs from clinical practice, and how a medicine travels from the laboratory to the market.

- Introduction to clinical research
- Clinical trial phases
- Pharmacological principles of clinical research
- Drug development and launch

MODULE 02 Regulations & Key Stakeholders

The rulebook and the people bound by it — who is accountable for what, and to whom.

- Indian regulation (domestic regulation)
- ICH GCP / Schedule Y / ICMR / Indian GCP
- Key stakeholders in clinical research
- Ethics Committees and Institutional Review Board
- Responsibilities of the sponsor and the investigator
- Sponsor–investigator and sponsor–vendor responsibilities

MODULE 03 Trial Design & Project Management

Designing a study that can actually answer its question, then running it to time and budget.

- Clinical trial design
- Vendor selection and management
- Project planning
- Project management

MODULE 04 GCP, Protocol & Essential Documents

The paperwork that is the trial — if it is not documented to GCP standard, it did not happen.

- Principles of Good Clinical Practice
- Protocol design and CRF design
- Essential documents and regulatory requirements
- IND application and NDA application
- Clinical study report
- Informed consent process and documentation

MODULE 05 Study Setup, Monitoring & Compliance

Opening a site, keeping it compliant and verifying at the source that the data are real.

- Study setup, site selection and pre-study visits
- Site initiation, subject recruitment and retention planning
- Site contracts and budgeting
- QC, compliance and auditing in clinical research
- 21 CFR Part 11 and site auditing
- Sponsor compliance, auditing and SOPs
- Clinical monitoring activities
- CRF review and source data verification
- Drug safety reporting and drug accountability
- Routine site monitoring and site close-out

MODULE 06 Foundations of Pharmacovigilance

Drug safety as a discipline — what it monitors, why it exists and the language it uses.

- Introduction to pharmacovigilance
- Overview of pharmacovigilance
- Standard terms and terminology in pharmacovigilance
- Global perspective of pharmacovigilance

MODULE 07 Adverse Events & Case Processing

From a single reported reaction to a properly assessed, coded and narrated safety case.

- Medical evaluation of adverse events
- Adverse event reporting system and form
- Diagnosis and management of ADRs
- Single case processing
- Case narrative writing

MODULE 08 Signal Detection, Risk & Aggregate Reporting

Moving from individual cases to a population-level view of a medicine's safety profile.

- PV reporting database and signal detection
- Signal management
- Risk assessment, evaluation and management
- Quality system in PV
- Expedited reporting criteria
- PSUR & PBRER

MODULE 09 MedDRA, PV Laws, Audits & Inspections

The terminology standard and the regulatory framework that govern every safety submission.

- Medical Dictionary for Regulatory Activities (MedDRA)
- Regulatory guidelines and laws in PV
- SOPs in PV
- PV auditing and inspection
- Regulatory aspects in PV

MODULE 10 · NEW Artificial Intelligence in Clinical Research & Pharmacovigilance

AI-Enabled Trials · Drug-Safety Intelligence · Responsible Human Oversight

- AI, Machine Learning, NLP, Generative AI and LLMs in life sciences
- AI across the clinical-trial lifecycle — planning, startup, conduct, monitoring, reporting
- AI-assisted protocol and clinical-study document review
- … and 19 further topics — the full module is set out on page 5

MODULE 10 · NEW HIGHLIGHT MODULE

Artificial Intelligence in Clinical Research & Pharmacovigilance

AI-Enabled Trials · Drug-Safety Intelligence · Responsible Human Oversight

This future-ready module introduces responsible applications of Artificial Intelligence, Machine Learning, Natural Language Processing and Generative AI across the clinical-research and pharmacovigilance lifecycle. The emphasis is practical AI literacy — understanding where AI can support professionals while participant protection, medical judgement, data integrity, confidentiality, traceability, validation and regulatory accountability are maintained.

Key Topics Covered

- AI, Machine Learning, NLP, Generative AI and LLMs in life sciences
- AI across the clinical-trial lifecycle — planning, startup, conduct, monitoring, reporting
- AI-assisted protocol and clinical-study document review
- AI support for literature review, evidence summarisation and knowledge management
- AI concepts for site feasibility, patient recruitment and retention
- AI-assisted clinical-data review, anomaly identification and quality signals
- AI in Risk-Based Quality Management (RBQM) and risk-based monitoring
- AI-assisted monitoring preparation, visit summaries and follow-up documentation
- AI support for essential-document review, checklists and inspection readiness
- AI-assisted adverse-event extraction from structured and unstructured sources
- AI concepts for safety-case intake, triage, duplicate detection and prioritisation
- AI-assisted case narrative drafting and quality review
- AI support for MedDRA coding suggestions and terminology mapping
- AI-assisted literature surveillance for drug-safety information
- AI concepts for signal detection, trend identification and safety-pattern recognition
- AI support for PSUR / PBRER organisation and consistency review
- Generative AI for SOPs, training material, audit questions and structured summaries
- Prompt engineering for clinical research and pharmacovigilance professionals
- AI hallucinations, fabricated citations, bias and inaccurate conclusions
- Participant privacy, confidential trial data and secure AI use
- Data integrity, audit trails, traceability, validation and 21 CFR Part 11
- Human-in-the-loop review and accountable medical / regulatory decision-making

THE RESPONSIBLE BOUNDARY

Anyone can generate an answer with AI. The professional skill — and the one employers are now paying for — is knowing how to find what the AI got wrong.

AI cannot replace qualified investigators, monitors, safety physicians, PV professionals or regulatory decision-makers. Every AI-generated clinical, safety or regulatory output must be compared with source data, approved documents and authoritative requirements before it is used.

What This Module Adds to Your Career

Clinical research and pharmacovigilance are document- and data-intensive professions, so AI literacy pays off directly in document review, data-quality signals, literature surveillance, case processing and structured drafting. Verification carries the emphasis, because AI-generated safety information can be confidently and completely wrong — and here an unverified statement is a patient-safety problem and a compliance problem at once.

AI-Enhanced Practice Exercises

Applied work that builds the verification habit, not just the prompting habit:

- Review an AI-generated protocol checklist for gaps and unsupported items
- Evaluate a draft case narrative against the source safety data
- Identify missing safety information in an AI-assisted case summary
- Detect hallucinated references in an AI literature summary
- Prepare a structured monitoring or PV quality checklist

HOW YOU STUDY & WHERE IT TAKES YOU

Study Method, Certification, Skills & Career Pathways

Study Methodology

- Freelance, self-paced method of study around your work schedule.
- Study the books and course material provided with your enrollment.
- Note down your queries as you progress through each module.
- Contact your assigned expert by email or phone at any time.
- Experts respond by mail, by call, or arrange a session based on your queries.
- Lifetime expert support to keep raising your knowledge and expertise.
- All content is based on live cases and industry best practice.

Mode of Examination

- Login details for the online examination are issued on completion of the programme duration.
- A 7-day window is given to appear for the online examination.
- The examination pattern is Multiple Choice Questions (MCQ).
- A PowerPoint project of minimum 30 pages on a topic of your own choice.
- Successful candidates are awarded two certifications (Degree and Transcriptional Degree).

Learning Mode, Assessment & Certification

Mode	Online and distance learning, designed for working professionals.
Assessment	Online and distance examinations (MCQ / theoretical) and/or projects.
Certification	GIHS certificate awarded to successful candidates on completion of applicable requirements.
Career support	Placement support and CV-forwarding assistance for successful candidates.
Verification	Employers can confirm certificate authenticity through the official GIHS verification process.

Verified by GIHS itself — not by a third party. Every certificate carries its own certificate ID. An employer, university or registration body can confirm it directly at www.gihsonline.com/verify-certification — no third-party verification agency and no fee.

Practical & Job-Oriented Learning

The programme connects concepts with real professional workflows: protocol and CRF review, informed-consent documentation, site setup, monitoring, source-data verification, drug accountability, adverse-event assessment, case processing, narrative writing, signal and risk-management thinking, SOPs, audits and inspection readiness. Learners practise the two habits the work depends on — checking a statement against its source, and documenting that the check happened.

Career-Relevant Skills You Will Develop

- Clinical trial lifecycle and GCP understanding
- Protocol, CRF and essential-document awareness
- Study setup and site-management concepts
- Clinical monitoring and source-data verification
- Quality, compliance and audit awareness
- Adverse-event and ADR terminology
- Case processing and case narrative concepts
- Signal detection and risk-management awareness
- PSUR / PBRER and MedDRA concepts
- PV quality systems, SOPs, audits and inspections
- AI literacy across clinical research and drug-safety workflows

Career & Professional Pathways

The current GIHS course page lists **CRA, Clinical Research Coordinator, Pharmacovigilance Associate and Drug Safety pathways**. The programme can also support:

- Clinical Research Associate (CRA)
- Clinical Research Coordinator (CRC)
- Clinical Trial Assistant / clinical operations support
- Pharmacovigilance Associate
- Drug Safety Associate / Drug Safety Specialist
- Clinical safety / PV operations support
- Site management and monitoring support
- Clinical research quality & compliance support
- PV quality / audit support
- Signal and safety surveillance support

Specific clinical, medical or senior roles may require additional qualifications, professional registration and relevant experience; GIHS professional education complements rather than replaces those requirements. GCP and drug-safety principles are internationally relevant, but candidates seeking work abroad must also meet destination-country employer, immigration and regulatory requirements.

PLACEMENT & CAREER ENHANCEMENT

What GIHS Does for Your Career After the Course

CAREER ENHANCEMENT INCLUDED FREE WITH EVERY ENROLMENT

Exclusive FREE CV & Interview Coaching — worth ₹9,999

FREE
worth ₹9,999

Join a GIHS programme now and unlock complimentary access to TopCV Coach, giving you the professional resume and interview skills you need to launch your career with confidence. Your profile is also posted on the TopCV Coach website, where a large recruiter and employer network is active.

Professional resume building
Build a CV that reads the way recruiters in healthcare actually screen.

Interview skills coaching
Practise and sharpen how you present your qualification and your new AI literacy.

Profile posted on TopCV
Your profile goes live on topcvcoach.com, in front of a large employer network.

No extra cost
Worth ₹9,999, included free with your GIHS enrolment.

TopCV Interview — Delivered with **www.topcvcoach.com** · included free with every GIHS programme enrolment

Where Our Graduates Are Making an Impact

Clinical research is a global industry, and GIHS learners work right across it — in pharmaceutical and biotech companies, CROs, trial sites, regulatory bodies, medical devices, healthcare IT and academic research.

Pharmaceutical companies
Clinical operations, drug safety and regulatory teams

Biotechnology & biosimilars
Development programmes and regulatory submissions

Contract Research Organisations
Monitoring, data management and safety services

Hospitals & trial sites
Coordination, documentation and ethics-committee support

Medical devices & diagnostics
Clinical evaluation, regulatory and quality functions

Healthcare IT & analytics
Clinical data platforms, SAS programming and reporting

Regulatory & consulting firms
Submissions, audits, IPR and compliance services

Academic & government research
Institutional trials, registries and regulatory bodies

Independent practice & ventures
Writing, data, PV and regulatory support services

WORKING GLOBALLY — India · UK · USA · Canada · Australia · New Zealand · UAE & Gulf · Saudi Arabia · Qatar · Oman · Russia · Germany · Ireland · Singapore · Malaysia · Africa · Europe

Placement support. More than 2,000 GIHS professionals are now working in well-known organisations worldwide, many of them after building CV and interview skills through the free TopCV Coach programme. GIHS shares your profile globally for career growth. Successful candidates receive placement support and their CVs may be forwarded to various organisations. Final employment remains subject to employer selection and role-specific eligibility.

Business & Entrepreneurship Value

Pharmacovigilance is a permanent obligation for every marketed product, so demand for safety capacity does not rise and fall with the trial pipeline — it accumulates.

- Provide pharmacovigilance case-processing and safety-operations support
- Offer literature-surveillance services to marketing-authorisation holders
- Build a clinical monitoring and site-management support practice
- Consult on PV quality systems, SOPs and audit readiness
- Deliver GCP and drug-safety training for pharma and CRO teams

Any regulated clinical service must only be provided within your lawful professional scope and applicable registration requirements. Career titles, eligibility and the right to practise depend on your prior qualification, experience, employer criteria and local regulation.

Global Institute Of Health Science

Certificate

OF POST GRADUATE DIPLOMA

This is to certify that

CANDIDATE NAME

has successfully completed the one year programme of **Post Graduate Diploma in Clinical Research & Pharmacovigilance** offered by Global Institute of Health Science, during **MONTH YEAR to MONTH YEAR**. Upon successful completion of the programme

POST GRADUATE DIPLOMA

IN

CLINICAL RESEARCH & PHARMACOVIGILANCE

is awarded to him/her this day the **XX MONTH YEAR**.

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Academic Head

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Global Institute Of Health Science

STATEMENT OF RESULT & STUDY

Grade Sheet & Transcript

CANDIDATE NAME	PROGRAMME PERIOD	TOTAL LEARNING HOURS
CANDIDATE NAME	MONTH YEAR – MONTH YEAR	600 Hours

PROGRAMME

Post Graduate Diploma in Clinical Research & Pharmacovigilance

NO.	MODULE & TOPICS COVERED	HOURS	MAX	OBTAINED	%
01	**Foundations of Clinical Research** Introduction to clinical research; Clinical trial phases; Pharmacological principles of clinical research; Drug development and launch	45	50	48.5	97%
02	**Regulations & Key Stakeholders** Indian regulation (domestic regulation); ICH GCP / Schedule Y / ICMR / Indian GCP; Key stakeholders in clinical research; Ethics Committees and Institutional Review Board; Responsibilities of the sponsor and the investigator; Sponsor–investigator and sponsor–vendor responsibilities	45	50	46.5	93%
03	**Trial Design & Project Management** Clinical trial design; Vendor selection and management; Project planning; Project management	45	50	47.5	95%
04	**GCP, Protocol & Essential Documents** Principles of Good Clinical Practice; Protocol design and CRF design; Essential documents and regulatory requirements; IND application and NDA application; Clinical study report; Informed consent process and documentation	45	50	47	94%
05	**Study Setup, Monitoring & Compliance** Study setup, site selection and pre-study visits; Site initiation, subject recruitment and retention planning; Site contracts and budgeting; QC, compliance and auditing in clinical research; 21 CFR Part 11 and site auditing; Sponsor compliance, auditing and SOPs; Clinical monitoring activities; CRF review and source data verification; Drug safety reporting and drug accountability; Routine site monitoring and site close-out	45	50	47	94%
06	**Foundations of Pharmacovigilance** Introduction to pharmacovigilance; Overview of pharmacovigilance; Standard terms and terminology in pharmacovigilance; Global perspective of pharmacovigilance	45	50	48	96%
07	**Adverse Events & Case Processing** Medical evaluation of adverse events; Adverse event reporting system and form; Diagnosis and management of ADRs; Single case processing; Case narrative writing	45	50	48	96%
08	**Signal Detection, Risk & Aggregate Reporting** PV reporting database and signal detection; Signal management; Risk assessment, evaluation and management; Quality system in PV; Expedited reporting criteria; PSUR & PBRER	45	50	49	98%
09	**MedDRA, PV Laws, Audits & Inspections** Medical Dictionary for Regulatory Activities (MedDRA); Regulatory guidelines and laws in PV; SOPs in PV; PV auditing and inspection; Regulatory aspects in PV	45	50	46	92%
10	**Artificial Intelligence in Clinical Research & Pharmacovigilance** NEW AI-Enabled Trials · Drug-Safety Intelligence · Responsible Human Oversight; AI, Machine Learning, NLP, Generative AI and LLMs in life sciences; AI across the clinical-trial lifecycle — planning, startup, conduct, monitoring, reporting; AI-assisted protocol and clinical-study document review	45	50	48.5	97%
	Module Examinations — Sub-total	**450**	**500**	476	**95.2%**
11	**Project Work** 30-page applied project submitted at the end of the programme	150	200	185	92.5%
	GRAND TOTAL	**600**	**700**	**661**	**94.4%**

TOTAL MARKS	PERCENTAGE	GRADE	RESULT	DATE OF AWARD	CERT. ID
661 / 700	94.4%	A+	PASSED	XX-XX-XXXX	gihs-XXXX-XXXXX

Academic Head

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ENROLMENT

Three Flexible Learning Options

Programme	Duration	Indian Fee	International	Positioning
Certificate Program	3 Months	**₹22,999**	**450 USD**	Basic foundation programme — core knowledge first
Post Graduate Diploma Program	1 Year	**₹32,999**	**699 USD**	Advanced and career-oriented — full depth at a steady pace
Fast Track PG Diploma Program MOST CHOSEN	4 Months	**₹36,999**	**750 USD**	Same advanced syllabus — flexible, finish at your own pace

Both advanced options carry the same career-oriented syllabus and the same two international certificates. The Fast Track is simply flexible — finish as quickly as your own pace allows and step into a fast-growing job market sooner. The Certificate Program is the basic, foundation-level option.

Printed books are optional
Online study material is included with every enrolment. Add the printed books at enrolment if you would like them shipped.

₹2,999 additional (India) | **99 USD** additional (International)

EMI available on credit cards Convert your fee into easy monthly instalments — ask admissions when you enrol.

All modes of payment are available. Fees may be subject to applicable taxes, exchange rates or future revision.

What Our Candidates Say

★★★★★ 300+ Google reviews

"I recently completed my Post Graduation in Pregnancy and Child Care from GIHS and it was a wonderful experience."
Dr Fatema Ebrahim

"One of the best institutions... Course is carefully structured and detailed. Management is very well organised and prompt in assistance."
A V

"This course has been a game-changer for me, providing the knowledge and skills necessary to succeed on a global platform."
Akram Sabiq

"They have a very good independent study methodology and the subject matter is really good... the Institute of Health was a wise choice."
Andrea Del Pilar

"From the day 1 to till date Chethan assisted very well... the studies are very detailed and made interesting."
Sarah Lilly, MBA

Admission Procedure START ANY DAY

1. **Apply online — any day**
   Enrol at www.gihsonline.com / Apply Online. Admissions stay open all year — no batch to wait for.
2. **Confirm your fee**
   No hidden charges. Payment within 7 days of enrollment.
3. **Receive your study material**
   Online access within 48 hours; printed books shipped if you add them.
4. **Get your personal expert**
   A professional subject expert is assigned to you for lifetime support at no cost.
5. **Complete exam and project**
   Online examination plus a project submission.
6. **Receive your certificates**
   Two international certificates are awarded and shipped.

Who Can Enrol

LISTED ON THE GIHS COURSE PAGE

- Science Graduates
- MBBS / BDS / BHMS / BAMS / BPT / BUMS
- B.Pharm / M.Pharm / Pharmacy Professionals
- Occupational Therapists
- B.Sc. / M.Sc. & Life-Science Graduates
- Medical IT Professionals
- Medical Laboratory Professionals
- Nursing Professionals
- Microbiology Professionals
- Biomedical / Biotechnology Professionals
- Clinical Research / CRO / Pharma Professionals
- Professionals working with hospitals & healthcare organisations

ALSO OPEN TO

- Any graduate from any discipline
- Working professionals with relevant work experience
- Entrepreneurs planning to start a healthcare or allied venture
- Managers and administrators running a healthcare organisation

Any graduate, and any professional with relevant work experience, may enrol — including those who intend to start, run or manage a healthcare or allied organisation.

QUESTIONS

Frequently Asked Questions

Why combine clinical research and pharmacovigilance?

The two are continuous. Trials generate safety information and pharmacovigilance provides the structured processes for evaluating and monitoring it, so learning both gives a broader view of product development and patient safety.

Does the programme cover ICH-GCP?

Yes. ICH-GCP and the principles of Good Clinical Practice are central to the syllabus, alongside Schedule Y, ICMR and Indian GCP.

Does it cover clinical monitoring?

Yes — clinical monitoring, CRF review, source-data verification, drug accountability, routine site monitoring and site close-out.

Does it cover case processing and narratives?

Yes. Case processing, single-case processing and case narrative writing are part of the pharmacovigilance curriculum.

Does it cover signal detection, PSUR/PBRER and MedDRA?

Yes. PV databases, signal detection and management, risk assessment, PSUR, PBRER and MedDRA are all specifically included.

Will AI replace a CRA or PV professional?

No. The module teaches AI as an assistive capability; clinical and safety work requires accountable human judgement, patient protection, data integrity and regulatory compliance.

Can this help me become a CRA or PV Associate?

It strengthens the knowledge these roles need. Actual appointment depends on prior qualification, competencies, experience and employer requirements.

Can working professionals enrol?

Yes. The programme is delivered online and by distance learning, with a Fast-Track option.

What is included in the programme fee?

Course material with the latest updates, online access within 48 hours, printed books shipped with your enrolment, a personal subject expert for life at no cost, the online examination and project assessment, and two international certificates on successful completion. There are no hidden charges.

Is the TopCV Coach access really free?

Yes. Every GIHS enrolment now includes complimentary TopCV Coach access worth ₹9,999 for professional resume building and interview skills, and your profile is posted on the TopCV website where a large recruiter network is active.

Can I use this programme to start my own practice or venture?

Yes, within your lawful professional scope. The Business & Entrepreneurship section of this brochure sets out the non-clinical services, education and consulting work the knowledge supports. Regulated clinical services require the appropriate qualification and authorisation.

I am not from a clinical background. Can I still enrol?

Yes. Any graduate from any discipline may enrol, as can working professionals with relevant experience, entrepreneurs and managers of healthcare or allied organisations.

No batch to wait for — start the day you enrol Admissions stay open every day of the year — enrol today and begin at once.

ABOUT THE INSTITUTE

Global Institute of Health Science

Global Institute of Health Science is a pioneer in health science education, especially in distance and online learning, with more than 15 years of experience and international exposure. GIHS is an ISO 21001:2018 certified health science institute, globally acclaimed, and certified with copyright approval from IPR, Government of India. GIHS also holds international certification from UASL UK for its Quality Control Management System.

- **Excellence for health professionals** — Programmes built around what healthcare employers actually ask for.
- **15+ years with international exposure** — A student and professional base across India, UK, USA, Canada, UAE, Russia and Australia.
- **ISO certified and government approved** — ISO 21001:2018, IPR Government of India copyright approval and UASL UK quality certification.
- **Lifetime free personalised tutoring** — A dedicated subject expert assigned to you at no cost, for life.

More Ways to Build Your Career

GIHS programmes help professionals get a strong start in the healthcare field. These courses are ad-hoc to a professional career and provide the additional benefit of being part of a global learning organisation.

Health Management Programs

- Health Informatics
- Hospital Administration
- Pharma Management
- Hospital Planning & Management
- Medical Transcription
- Nursing Management
- Public Health Administration
- Health Insurance
- Blood Banking Management
- Medico-Legal Systems
- Medical Laboratory Technology
- Medical Radiology Technology
- Quality Management for Hospitals
- Disaster Management for Hospitals
- Hospital Infection Control & Patient Safety
- Healthcare & Hospital Administration
- Medical Record Management
- Medical Scribe

Clinical Research Programs

- Clinical Research & Quality Assurance
- Pharmacovigilance & Medical Writing
- Clinical Research & Pharmacovigilance
- Clinical Data Management & SAS
- Regulatory Affairs, IPR & Patent
- Clinical Trials Management
- CRA & Site Management
- Study Start-Up & Site Activation
- Trial Supply & IP Management
- Biostatistics & Statistical Programming

Allied Medical Programs

- Mother & Child Health
- Mental Health & Addictions
- Nutrition & Dietetics
- Pregnancy Education
- Gynaecology & Obstetrics
- Electroencephalogram (EEG)
- Nuclear Medicine Technology
- Anaesthesia Management
- First Aid (CPR / Emergency Care)
- Electrocardiogram (ECG)

Environment, Health & Safety Programs

- Industrial Health & Hygiene
- Environment Health Safety & RM
- Occupational Health & Safety
- Industrial Safety
- Food Safety & Quality Management
- Oil & Gas Safety Engineering
- Process Safety & Engineering
- Fire & Industrial Safety Management
- Environment Social Governance (ESG)

Ayurveda & Homeopathic Programs

- Panchkarma: Ayurveda
- Garbhsanskar: Ayurveda
- Naturopathy & Yoga
- Siddha: Medicine
- Ksharsutra: Ayurveda
- Classical Homeopathy
- Materia Medica: Homeopathy
- Repertory: Homeopathy
- Organon: Homeopathy
- Ayurvedic Nutrition & Dietetics

Ready to enrol?

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